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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/571,012	03/08/2006	Frank Cuttitta	4239-82094-06	4600
36218 7590 01/05/2011 KLARQUIST SPARKMAN, LLP (OTT-NIH) 121 S.W. SALMON STREET SUITE #1600 PORTLAND, OR 97204-2988			EXAMINER	
			PAGONAKIS, ANNA	
			ART UNIT	PAPER NUMBER
			1628	
			NOTIFICATION DATE	DELIVERY MODE
			01/05/2011	ELECTRONIC

# Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

tanya.harding@klarquist.com docketing@klarquist.com

	Application No.	Applicant(s)				
Office Action Cumment	10/571,012	CUTTITTA ET AL.				
Office Action Summary	Examiner	Art Unit				
	ANNA PAGONAKIS	1628				
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	correspondence address				
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.  - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.  - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.  - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).						
Status						
1) Responsive to communication(s) filed on 20 De	ecember 2010.					
2a) ☐ This action is <b>FINAL</b> . 2b) ☐ This action is non-final.						
3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is						
closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.						
Disposition of Claims						
4) Claim(s) 80,81,90-97 and 100-103 is/are pending in the application.						
4a) Of the above claim(s) <u>91-93</u> is/are withdrawn from consideration.						
5) Claim(s) is/are allowed.						
6) Claim(s) <u>80,81,90,94-97 and 100-103</u> is/are re	jected.					
7) Claim(s) is/are objected to.						
8) Claim(s) are subject to restriction and/or election requirement.						
Application Papers						
9) The specification is objected to by the Examiner.						
10) ☐ The drawing(s) filed on is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.						
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).						
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).						
11) $\square$ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.						
Priority under 35 U.S.C. § 119						
12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).						
a) All b) Some * c) None of:						
1. Certified copies of the priority documents have been received.						
2. Certified copies of the priority documents have been received in Application No						
3. Copies of the certified copies of the priority documents have been received in this National Stage						
application from the International Bureau (PCT Rule 17.2(a)).						
* See the attached detailed Office action for a list	of the certified copies not receive	ed.				
Attachment(s)						
1) Notice of References Cited (PTO-892)  4) Interview Summary (PTO-413) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948)  Paper No(s)/Mail Date						
2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08)	3) Notice of Dransperson's Patent Drawing Review (PTO-946)  Information Disclosure Statement(s) (PTO/SB/08)  Notice of Informal Patent Application					
Paper No(s)/Mail Date 1 sheet, 12/20/2010	6) Other:					
U.S. Patent and Trademark Office PTOL-326 (Rev. 08-06) Office Ac	tion Summary Pa	art of Paper No./Mail Date 20101229				

### **DETAILED ACTION**

Applicant's amendment filed 12/20/2010 have been received and entered into the present application.

As reflected by the attached, completed copy form PTO/SB/08A (one page total), the Examiner has considered the cited references.

Applicant's arguments filed 12/20/2010 have been fully considered. Rejections not reiterated from previous Office Actions are hereby withdrawn. The following rejections are either reiterated or newly applied. They constitute the complete set of rejections presently being applied to the instant application.

#### Status of Claims

Claims 80-81, 90-97 and 100-103 are pending.

Claims 91-93 are withdrawn.

Claims 80-81, 90, 94-97 and 100-103 are currently under examination and the subject matter of the present Office Action.

## Rejection maintained:

### Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

Claims 80-81, 90, 94-97 and 100-103 are rejected under 35 U.S.C. 102(b) as being anticipated by JP 10212235 (original document and translation are attached) as evidenced by the National Cancer Institute (1/28/2005).

Art Unit: 1628

JP 10212235 teaches a compound of formula (I) wherein the compound is that of instant claim 80 (paragraph [0064] of translation):

Further, it is taught that the compounds of formula (I) is effective for the treatment of tumors, for example, stomach cancer, such as malignant tumor, a benign tumor, and a precancerous change, lung cancer, hepatoma, a pancreatic cancer, colon cancer, a malignant lymphoma, leukemia, a breast carcinoma, melanoma, renal cancer, brain tumor, peritoneal tumor, spinal cord tumor, hypophyseal tumor, thyroid tumor, laryngeal cancer etc. (paragraph [0035] of translation).

Though JP 10212235 is silent as to the effect of the elected compound to inhibit an activity of a gastrin releasing peptide (GRP), the administration of the claimed compound to patients suffering from cellular proliferative disorders is expected to necessarily have the claimed effect of inhibiting an activity of GRP, whether recognized by the author or not. Products of identical chemical composition cannot exert mutually exclusive properties when administered under the same circumstance or, in the present case, the same host. Please reference MPEP 2112.

Moreover, the very teaching of administering the identical compound to the same patient populations (i.e. patients suffering from cellular proliferative disorders) necessarily means that the claimed inhibition of GRP is necessarily present, whether recognized by the author or not. Further, the

Art Unit: 1628

treatment of a tumors would necessarily inhibit angiogenesis since angiogenesis is responsible for the progression of the disease (i.e. is angiogenesis mediated), per National Cancer Institute. As stated supra, products of identical composition cannot exert mutually exclusive properties. Please reference MPEP 2112 and Ex parte Novitski, 26 USPQ 1389 (Bd. Pat. App. and Inter 1993).

#### Response to Applicant's Remarks

Applicant alleges that JP10212235 does not describe the effects of any compound on GRP activity, nor is GRP mentioned in the document. This is not found persuasive. Though JP 10212235 is silent as to the effect of the elected compound to inhibit an activity of a gastrin releasing peptide (GRP), the administration of the claimed compound to patients suffering from cellular proliferative disorders is expected to necessarily have the claimed effect of inhibiting an activity of GRP, whether recognized by the author or not. Products of identical chemical composition cannot exert mutually exclusive properties when administered under the same circumstance or, in the present case, the same host. Please reference MPEP 2112.

Applicant alleges that "without any admission to any similarity between Compound 105 and claimed Compound 77427", compound 105 is not among the compounds used in vitro tests of anti-proliferative activity. Firstly, Applicant's statement is not clear. If Applicant does not believe that compound 105 of JP 10212235 is the instantly elected compound, Applicant is invited to set forth reasons and evidence. Further, JP 10212235 clearly states that compounds of formula (I), including the elected compound, are useful for the treatment of tumors, for example, stomach cancer, such as malignant tumor, a benign tumor, and a precancerous change, lung cancer, hepatoma, a pancreatic cancer, colon cancer, a malignant lymphoma, leukemia, a breast carcinoma, melanoma, renal cancer, brain tumor, peritoneal tumor, spinal cord tumor, hypophyseal tumor, thyroid tumor, laryngeal cancer etc. (paragraph [0035] of translation).

Applicant alleges that Tables 27-31 does not show an anti-proliferative effect on every compound against every cell line tested. This is not found persuasive. It appears from this statement that Applicant infers requirement of testing "every cell line." Applicant is reminded that as stated in the Final Rejection mailed on 6/10/2010 the instant specification merely teaches proliferation assay of lung cancer cell line H1299 with administration of the elected compound as well as xenograft lung cancer model (Examples 8 and 9, pages 36-37). Therefore, it seems that Applicant is arguing against the enablement of their own invention.

Applicant alleges that the omissions in some of the tables demonstrate considerable variability among the biological effects of the tested compounds. It is not clear how Applicant has reached this conclusion. Applicant fails to advanced any specific reasons or evidence, aside from Counsel's own allegation, in support of this position that no motivation exists in the present obviousness rejection. This assertion by Counsel is an unsupported allegation and fails to take the place of evidence in the record. Statements of this nature are clearly unpersuasive in accordance with the guidance provided at MPEP 2145, which states "The arguments of counsel cannot take the place of evidence in the record."

Applicant alleges that the compound used in in vitro and in vivo experiments differ structurally from the instantly claimed compound, as presented in Exhibit A. It is not clear what Exhibit Applicant is referring to. The only Exhibit A previously submitted was a Curriculum Vitae of Frank Cuttitta filed on 9/10/2010.

#### Conclusion

No claim is found to be allowable.

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to ANNA PAGONAKIS whose telephone number is (571)270-3505. The examiner can normally be reached on Monday thru Thursday, 7am to 5pm EST.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Brandon Fetterolf can be reached on 571-272-2919. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Application/Control Number: 10/571,012

Page 7

Art Unit: 1628

Supervisory Patent Examiner, Art Unit 1628